

Dated: September 23, 2019.
Thomas W. Prevoznik,
Acting Assistant Administrator, Deputy Assistant Administrator.
 [FR Doc. 2019–21320 Filed 9–30–19; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: The registrants listed below have applied for and been granted a

registration by the Drug Enforcement Administration (DEA) as a bulk manufacturer of a various classes of schedule I and II controlled substances.
SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as a bulk manufacturer of various classes of scheduled I and II controlled substances. Information on a previously published notices is listed below. No comments or objections were submitted for these notices.

Company	FR docket	Published
Absolute Standards, Inc	84 FR 31620	July 2, 2019.
Pisgah Laboratories, Inc	84 FR 31622	July 2, 2019.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing each company’s physical security systems, verifying each company’s compliance with state and local laws, and reviewing the company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a

registration as a bulk manufacturer to the above listed companies.

Dated: September 24, 2019.
Thomas W. Prevoznik,
Acting Assistant Administrator, Deputy Assistant Administrator.
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DEPARTMENT OF JUSTICE
Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration; Siemens Healthcare Diagnostics, Inc.

ACTION: Notice of registration.

SUMMARY: The registrants listed below have applied for and been granted a registration by the Drug Enforcement Administration (DEA) as a bulk manufacturer of various classes of schedule I and II controlled substances.
SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as a bulk manufacturer of a basic class of schedule I and II controlled substances. Information on previously published notices is listed below. No comments or objections were submitted for the notice.

Company	FR Docket	Published
Siegfried USA, LLC	84 FR 7129	March 1, 2019.
Patheon Pharmaceuticals, Inc	84 FR 8114	March 6, 2019.
S & B Pharma Inc	84 FR 8116	March 6, 2019.
Siemens Healthcare Diagnostics, Inc	84 FR 10534	March 21, 2019.
Synthcon, LLC	84 FR 13962	April 8, 2019.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company’s maintenance of effective controls against diversion by inspecting and testing each company’s physical security systems, verifying each of the company’s compliance with state and local laws, and reviewing each company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR

1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed companies.

Dated: September 23, 2019.
Thomas W. Prevoznik,
Acting Assistant Administrator, Deputy Assistant Administrator.
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DEPARTMENT OF JUSTICE
Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Noramco, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 31, 2019. Such persons may also file a written request for a hearing on the application on or before October 31, 2019.